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(21) International Application Number: PCT/IB97/01253 (22) International Filing Date: 9 October 1997 (09.10.97) (30) Priority Data: <table border="0"><tr><td>08/729,933</td><td>15 October 1996 (15.10.96)</td><td>US</td></tr><tr><td>08/730,136</td><td>15 October 1996 (15.10.96)</td><td>US</td></tr><tr><td>08/730,554</td><td>15 October 1996 (15.10.96)</td><td>US</td></tr></table> (71) Applicant: RHONE-POULENC INC. [US/US]; Prospect Plains Road, Cranbury, NJ 08512-7500 (US). (72) Inventor: CUIF, Jean, Pierre; 42 Markham Road, Princeton, NJ 08540 (US). (74) Agents: SHEDDEN, John, A. et al.; Rhone-Poulenc Inc., Prospect Plains Road, Cranbury, NJ 08512-7500 (US).		08/729,933	15 October 1996 (15.10.96)	US	08/730,136	15 October 1996 (15.10.96)	US	08/730,554	15 October 1996 (15.10.96)	US	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
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(54) Title: CERIUM AND ZIRCONIUM OXIDES, MIXED OXIDES AND SOLID SOLUTIONS HAVING IMPROVED THERMAL STABILITY FOR CATALYSIS OF EXHAUST GAS SYSTEMS AND METHODS OF PRODUCING											
(57) Abstract <p>Mixed oxides of cerium and zirconium as well as solid solutions and cerium oxides are produced having improved thermal stability. The process involves the formation of cerium and zirconium oxides, mixed oxides and solid solutions by methods such as co-thermohydrolysis and co-precipitation, followed by washing or impregnating with an alkoxyated compound having greater than (2) carbon atoms. After the washing or impregnating step, the mixture is calcined to form oxides having very high surface area and oxygen storage capacity. The improved thermal stability of the oxides makes them advantageous for use in applications such as catalytic converters.</p>											

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**CERIUM AND ZIRCONIUM OXIDES, MIXED OXIDES AND SOLID
SOLUTIONS HAVING IMPROVED THERMAL STABILITY FOR
CATALYSIS OF EXHAUST GAS SYSTEMS AND METHODS OF
PRODUCING**

20

BACKGROUND OF THE INVENTION

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Field of the Invention

The present invention relates particularly to the production of mixed oxides, and more particularly relates to the production of cerium and zirconium mixed oxides having improved thermal stability. The mixed oxides have fine particle size distribution, very high surface area, oxygen storage and release capacity, and are useful in many applications including catalytic converters.

Background Information

Mixed oxides of cerium and zirconium are used for many applications, including catalysts used in automotive catalytic converters and the like. Such cerium and zirconium mixed oxides are typically formed by known precipitation techniques which involve the formation of the solid oxides in a liquid medium. When such mixed oxides are to be used, for example, in catalytic converters, it is desirable to maximize the thermal stability of the compounds, as defined by the stability of the surface area of the material after aging at high temperature. It is also desirable to maximize the surface area of such mixed oxides in order to provide improved catalytic properties. In addition to mixed oxides the present invention also relates to cerium oxides, zirconium oxides, and cerium/zirconia solid solutions (where substitution between cerium and zirconium in the network of the oxide as opposed to being two different phases is one phase).

Most industrial processes which include the precipitation or the creation of a solid in a liquid medium involve a solid/liquid separation state. Filtration, decantation or centrifugation are among the known techniques used for this purpose. After the solid/liquid separation is completed, the so-called wet cake
5 comprises precipitated particles and remaining mother liquor. In most of the processes, the mother liquor contains some salts that can contaminate the oxides generated during the next calcination operation. To reduce the amount of contaminants, washing is needed after the solid/liquid separation. In cases where the salts used as raw materials to make the precipitation are soluble in water,
10 washing is typically carried out with water. The volume and temperature of water used for washing determine the purity of the material and its thermal stability as well.

Although such techniques have been used to produce cerium and zirconium mixed oxides, a need still exists for the production of such mixed oxides
15 and solid solutions having substantially improved thermal stability. It is an object of the present invention to provide mixed oxides of improved thermal stability as well as cerium and zirconium oxides, preferably cerium oxides of improved thermal stability in addition to solid solutions having improved thermal stability.

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SUMMARY OF THE INVENTION

One aspect of the invention is to provide a novel way to improve the thermal stability of cerium and zirconium oxides, $(\text{Ce,Zr})\text{O}_2$ mixed oxides, solid solutions, and mixtures of cerium oxides and zirconium oxides, obtained by processes such as co-precipitation or thermohydrolysis, by introducing alkoxylated
25 compounds by impregnating or in the washing solution of a filter cake during the solid/liquid separation. The cake resulting from the precipitation typically occurs in aqueous media. By washing or impregnating with an alkoxylated compound, thermal stability can be substantially improved. Impregnating involves not draining the cake and instead making a paste which is contacted with the alkoxylated
30 compound.

An object of the present invention is to provide a method for producing cerium and zirconium oxides and mixed oxides, solid solutions and mixtures of oxides having improved thermal stability. The method is preferred for use in producing cerium oxides, cerium and zirconium mixed oxides and cerium and zirconium solid solutions having improved thermal stability. The method includes the steps of forming a mixture comprising a cerium and zirconium mixed oxide ((Ce, Zr)O₂) or a (Ce,Zr)O₂ solid solution or a cerium oxide (CeO₂) or zirconium oxide (ZrO₂), washing or impregnating with an alkoxylated compound, and drying or calcining the washed product. In a preferred embodiment, the product can be washed with water prior to washing with the alkoxylated compound. The alkoxylated compound is preferably provided in the form of an aqueous solution.

Another object of the present invention is to provide cerium and zirconium oxides, mixed oxides, mixtures of oxides and solid solutions having improved thermal stability. The oxides, mixed oxides and solid solutions have very high surface areas, very high oxygen storage capacities and low particle size.

These and other objects of the present invention will be more readily apparent from the following description.

All ratios, proportions and percentages herein are by weight, unless otherwise specified. "Comprising," as used herein, means various components can be conjointly employed. Accordingly, the terms "consisting essentially of" and "consisting of" are embodied in the term "comprising."

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The thermal stability of inorganic compounds can be defined as the stability of the surface area when material is aged at high temperature. For many applications, particularly catalysis, high surface area and highly stable materials are required by end users. In accordance with the present invention, cerium and zirconium mixed oxides and solid solutions are produced having improved thermal stability. The invention is also useful for producing cerium oxides, zirconium oxides and mixtures of cerium oxides and zirconium oxides having improved thermal stability.

The present invention relates to the use of alkoxyated compounds having greater than 2 carbon atoms during the washing or impregnating stage in order to improve the thermal stability of (Ce,Zr)O₂ mixed oxides and solid solutions. The alkoxyated compounds suitable for use herein have greater than 2 carbon atoms. The mixed oxides and solid solutions can be prepared by various conventional processes. Salts of Ce(III) and Zr(IV), nitrates for instance, can be mixed together and precipitated out by adding a base such as sodium hydroxide or ammonia. Adequate precipitation conditions must be used to obtain the mixed oxide phase after calcination at high temperature. Another process can involve the use of Ce(IV) as a starting material together with a Zr salt. This process also requires the use of a base as a precipitating agent. In any case, the precipitate has to be separated from the mother liquor. In the scope of the present invention, the solid is separated from the liquid by filtration or any other suitable method. In a preferred embodiment, the solid, otherwise called wet cake, is washed during a first stage with water to remove the water-soluble salts, nitrates for instance if nitrate solutions are the raw materials for the reaction. In a second stage of the preferred embodiment, the wet cake is washed or impregnated with a solution containing alkoxyated compounds such as ethoxylated alcohols, organic compounds or ethoxylated polymers such as PEG. Once washed or impregnated, the wet cake is either dried and calcined or directly calcined. The final product is a pure mixed oxide having substantially no organics since they are burned during calcination.

The (Ce,Zr)O₂ mixed oxides are initially formed by conventional processes such as co-thermohydrolysis or co-precipitation. Each of these processes is generally described separately below.

Co-thermohydrolysis

The first stage of the co-thermohydrolysis process involves preparing a mixture, in aqueous medium, of at least a soluble cerium compound and at least a soluble zirconium compound. The mixture can be obtained either from solid compounds which are dissolved in water, or directly from aqueous solutions of these compounds, followed by mixing, in any order, of the defined solutions.

Of the water soluble cerium compounds, one example is Ce IV salts, such as nitrates including ceric ammonium nitrate, that are suitable for the present invention. Preferably, a cerium nitrate is used. The cerium IV salt solution can contain some cerium III. However, it is preferred that the salt contains at least
5 about 85% cerium IV. An aqueous solution of cerium nitrate can be obtained by reacting nitric acid with a hydrated ceric oxide, prepared by a standard reaction of cerium III salt solution, carbonate for instance, with an ammonia solution in the presence of hydrogen peroxide, an oxidizing agent.

The aqueous solution of cerium IV salt can have some free acid, for
10 instance a normality ranging from 0.1 to 4 N. In the present invention, it is possible to use either a solution containing some free acid or a pre-neutralized solution by addition of a base, such as an aqueous solution of ammonia or alkaline hydroxides, e.g., sodium, potassium, etc. Preferably an ammonia solution is used to reduce the free acidity. In this case, it is possible to define the neutralization rate (r) of the
15 initial solution by the following equation:

$$r = (n_3 - n_2) / n_1$$

wherein n_1 represents the total number of moles of Ce IV present in the solution
20 after neutralization, n_2 represents the number of OH^- ions effectively used to neutralize the initial free acidity from the Ce IV aqueous solution, and n_3 represents the total number of moles of OH^- ions from the base added. When a neutralization step is used, excess base can be used in order to ensure the complete precipitation of the $\text{Ce}(\text{OH})_4$ species. Preferably, r is lower than 1, more preferably about 0.5.

25 The soluble zirconium salts used in the invention can be, for instance, zirconium sulfate, zirconyl nitrate or zirconyl chloride.

The amount of cerium and zirconium contained in the mixture substantially corresponds to the stoichiometric proportion required to obtain the final desired composition.

Once the mixture is obtained, it is then heated. The thermal treatment, called thermohydrolysis, is carried out at a preferred temperature of between about 80°C and the critical temperature of the reacting medium, typically between about 80 and about 350°C, more preferably between about 90 and about 5 200°C.

The heating stage can be carried out under air or under an inert gas such as nitrogen. Any suitable reaction time can be used, usually between about 2 and about 24 hours. The thermal treatment can be performed under atmospheric pressure or under any higher pressure such as the saturated vapor pressure. When 10 the temperature is higher than the reflux temperature of the reaction medium (usually higher than about 100°C), for instance between about 150 and about 350°C, the reaction is performed in a closed reactor or autoclave. The pressure can be equal to the autogenic pressure and can be correlated to the chosen temperature. It is also possible to increase the pressure in the reactor.

15 After the heating stage, a solid precipitate is recovered from the reactor and separated from the mother liquor by any process known by the state of art, for example filtration, settling or centrifugation.

If required, some additional base can be added directly after the heating stage into the reaction medium in order to improve the yield of the reaction. 20 The obtained precipitate can be washed or impregnated with one or several alkoxylated compounds, as more fully described below. In one embodiment, the precipitate is then dried, under air conditions for instance, at a temperature ranging from about 80 to about 300°C, preferably from about 100 to about 150°C. The drying stage is preferably performed until substantially no more weight loss is 25 observed.

After the optional drying step, the recovered precipitate is then calcined. This allows the formation of a crystalline solid solution phase. Usually, the calcination is carried out at temperatures ranging from about 200 to about 1000°C. The calcination temperature is typically higher than about 300°C, and 30 preferably ranges from about 400 to about 800°C.

Co-precipitation

The first stage of the co-precipitation process is the preparation of a mixture in an aqueous medium of at least a soluble cerium compound and at least a soluble zirconium compound. The mixture can be obtained either from solid compounds which are dissolved in water, or directly from aqueous solutions of these compounds, followed by mixing, in any order, of the defined solutions.

Suitable water soluble cerium compounds include cerium III salts, like nitrates or chlorides, for instance.

The soluble zirconium salts used in the invention can be, for instance, zirconium sulfate, zirconyl nitrate or zirconyl chloride.

The amount of cerium and zirconium contained in the mixture must correspond to the stoichiometric proportion required to obtain the final desired composition. Hydrogen peroxide can be added to the cerium/zirconium mixture or to the cerium or zirconium salt before mixing together. The effect of hydrogen peroxide is to oxidize at least part of Ce III to Ce IV.

Once the mixture is prepared, a base can be added to the cerium/zirconium salt solution to precipitate out the hydroxides. The base can be an ammonia solution or alkaline hydroxide solution, e.g., sodium, potassium, etc. The base solution used can, in particular, be an aqueous solution of ammonia or of sodium or potassium hydroxide. Ammonia solution is preferably used.

The precipitation is carried out on a batch or continuous basis. In the case of a continuous precipitation, the pH of the reaction is typically maintained between about 7 and about 11, preferably between about 7.5 and about 9.5. The residence time of the material in the reactor is typically at least about 15 minutes, preferably about 30 minutes. The reaction can be carried out at any suitable temperature such as room temperature. In the case of batch precipitation, the amount of base added is preferably at least the amount required to precipitate out $\text{Ce}(\text{OH})_4$ and $\text{Zr}(\text{OH})_4$.

After the reaction stage, a solid precipitate is recovered from the reactor and separated from the mother liquor by any process known by the state of art, for example filtration, settling or centrifugation. The obtained precipitate can then be washed or impregnated with one or several alkoxyated compounds, as
5 described below.

The next stage of the process is calcination of the washed material, either with or without an intermediate drying step. This allows the formation of a crystalline solid solution phase. Usually, the calcination is carried out at temperatures ranging from about 200 to about 1000°C. Calcination temperatures of
10 greater than about 300°C are suitable, preferably ranging from about 350 to about 800°C.

The alkoxyated compounds of the present invention can be defined by the general formula:



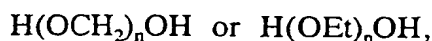
wherein R_1 and R_2 represent linear or non-linear alkyl groups or H or OH or Cl or Br or I; n is a number from 1 to 100; and x is a number from 1 to 4. R_1 and R_2 can contain an alcohol group. Of the alkyl groups, methoxy, ethoxy and propoxy
20 groups are preferred in order to generate an improvement in the thermal stability of the (Ce,Zr)O₂ mixed oxides and solid solutions.

The alkoxyated compound can be of the formula:



wherein R_1 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons and fatty hydrocarbon residues having from 8 to 20 carbons, n is from 1 to 100, and x is from 1 to 4. Preferably, n is from 12 to 40 and x is from 1 to 3. More preferably, x is 2.

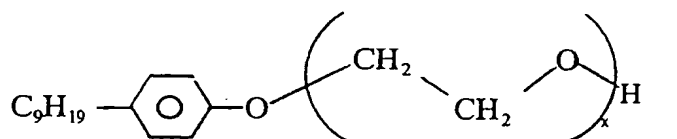
The alkoxyated compound can be of the formula:



wherein the average of n is from 1 to 100.

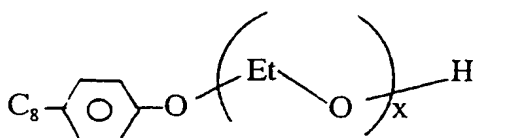
Examples of suitable alkoxyated compounds can be of the formulae:

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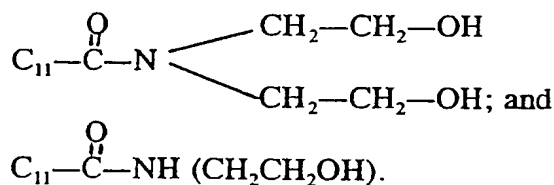
wherein the average of x is 9 or from 4 to 15;

10



wherein the average of x is 12.5;

15



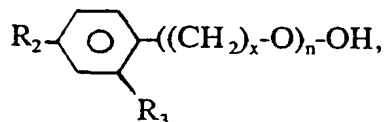
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Commercially available alkoxyated compounds suitable for use are sold by Rhône-Poulenc Inc. under the tradenames: IGEPAL CO 630, IGEPAL CA 720, ALKAMIDE LE, and ALKAMIDE L203.

25

The alkoxyated compound can alternatively be of the formula:

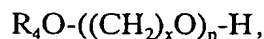
30



wherein R_2 and R_3 are the same or different and are independently selected from the group consisting of hydrogen and linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4. Preferably, n is from 12 to 40 and x is from 1 to 3. More preferably, x is 2.

35

The alkoxylated compound can further be of the formula:



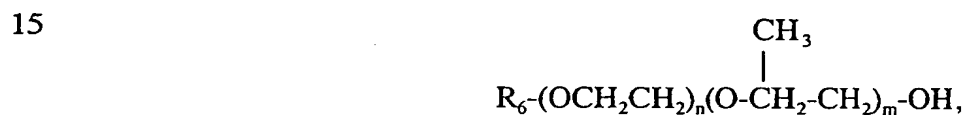
- 5 wherein R_4 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4. Preferably, n is from 12 to 40 and x is from 1 to 3. More preferably, n is 3, and x is 2.

The alkoxylated compound can further be of the formula:



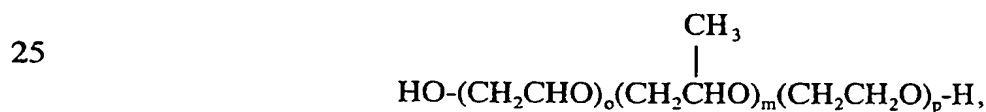
- 10 wherein R_5 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4. Preferably, n is from 4 to 40 and x is from 1 to 3. More preferably, x is 2.

The alkoxylated compound can also be of the formula:



- wherein R_6 is selected from the group consisting of linear and nonlinear alkyl groups
20 having from 1 to 20 carbons, n is from 1 to 100, and m is from 0 to 300 preferably 0 to 100. Preferably, n is from 12 to 40 and m is from 1 to 40.

The alkoxylated compound can alternatively be of the formula:



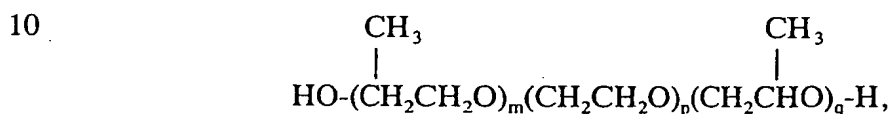
wherein o is from 0 to 300, m is from 0 to 300, and p is from 0 to 300.

The alkoxyated compound can further be of the formula:



- 5 wherein R_7 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4. Preferably, n is from 4 to 40 and x is from 1 to 3. More preferably, x is 2.

The alkoxyated compound can further be of the formula:



- 15 wherein m is from 0 to 300, p is from 0 to 300, and q is from 0 to 300 and having an average molecular weight of from about 40 to about 8,000.

In accordance with the present invention, the alkoxyated compound can comprise nonionic compounds. For example, the alkoxyated compound can comprise, or be derived from, compounds as listed below:

- 20 - Polyoxyalkylenated (polyethoxyethylenated, polyoxypropylenated, polyoxybutylenated) alkylphenols in which the alkyl substituent is C_6 - C_{12} and containing from 5 to 25 oxyalkylene units; examples include Tritons X-45, X-114, X-100 and X-102, marketed by Rohm & Haas Co.;
- 25 - glucosamide, glucamide, and glycerolamide;
- 30 - polyoxyalkylenated C_6 - C_{22} aliphatic alcohols containing from 1 to 25 oxyalkylene (oxyethylene, oxypropylene) units; examples include Tergitol 15-S-9 and Tergitol 24-L-6, marketed by Union Carbide Corp.; Neodol 45-9, Neodol 23-65, Neodol 45-7 and Neodol 45-4, marketed by Shell Chemical Co.; and Kyro KOB marketed by the Procter & Gamble Co.;

- 5 - the products resulting from the condensation of ethylene oxide, the compound resulting from the condensation of propylene oxide with propylene glycol, such as the Pluronics marketed by BASF;
- the products resulting from the condensation of ethylene oxide, the compound resulting from the condensation of propylene oxide with ethylenediamine, such as the Tetronics marketed by BASF;
- 10 - amine oxides such as (C₁₀-C₁₈ alkyl) dimethylamine oxides and (C₆-C₂₂ alkoxy) ethyldihydroxyethylamine oxides;
- the alkylpolyglycosides described in U.S. Patent No. 4,565,647;
- C₆-C₂₀ fatty acid amides;
- C₆-C₂₀ alkamides, preferably utilized at low concentrations;
- 15 - ethoxylated fatty acids; and
- ethoxylated amines.

 The alkoxyated compound of the present invention can advantageously be provided in the form of an aqueous solution having a relatively minor amount of the alkoxyated compound. The alkoxyated compound preferably
20 comprises less than about 50 wt.% of the aqueous solution, and more preferably comprises from about 0.1 to about 30 wt.% of the aqueous solution. A preferred commercially available compound suitable for use is sold by Rhône-Poulenc Inc. under the tradename IGEPAL 41.

 The final calcined mixture produced in accordance with the present
25 invention typically has a weight ratio of CeO₂ to ZrO₂ of from about 0:100 to about 100:0, preferably from about 95:5 to about 5:95. The resultant calcined mixture has a very high surface area, for example, greater than about 50m²/g. The final calcined mixed oxides also have a very high oxygen storage capacity, for example, greater than about 3ml O₂/g.

The surface area of the mixed oxides produced in accordance with the present invention is designated as B.E.T. determined by nitrogen adsorption according to the standard procedure ASTM D 3663-78 established from the method by BRUNAUER -EMMET - TELLER described in Journal of the American
 5 Chemical Society, 60, 309 (1938). Thermal stability is designated as the surface area of any powdery inorganic material after aging at a given temperature for a certain time. In the current invention, 10 g of material are calcined in a muffle furnace for about 6 hours at about 900°C. After this aging stage, the surface area of the material is measured by the method previously described.

10 The following examples illustrate various aspects of the invention and are not intended to limit the scope thereof.

Example 1

Using a co-precipitation method from a nitrate solution of cerium and zirconium, a mixed oxide is prepared of the composition: 75 wt% CeO₂ and 25
 15 wt% ZrO₂. The prepared nitrate solution comprises:

Ce(NO ₃) ₃ *	2856.24 g,
ZrO(NO ₃) ₂ *	2097.96 g,
H ₂ O ₂ (30%H ₂ O ₂)	568.92 g and
Deionized Water	8205.60 g;

20 * Rare Earth Lanthanide Series impurities such as Nd(NO₃)₃, Pr(NO₃)₃, and/or La(NO₃)₃ can be present in an amount of about 124 g.
 with the components being mixed well until homogenous.

The prepared ammonia solution comprises:

Ammonia solution (29.8% NH ₃)	- 1394.4 g, and
25 Deionized water	- 10365.6 g;

with the components being mixed well until homogenous.

The nitrate solution is added to the ammonium solution to precipitate out the hydroxides. The pH of the reaction is maintained between about 7 and 11. The temperature is room temperature. By reacting the appropriate salts and adding
 30 ammonia, the mixed hydroxides, corresponding to about 30 g of dried rare earth

oxides, are precipitated out from the solution and filtered on a Buchner filter. The cake is washed with about 750 ml of deionized water, then calcined for about 2 hours at about 400°C. The thermal stability of the product is evaluated after calcination in a muffle furnace at about 900°C for about 6 hours, and the surface area is measured using the B.E.T. method. The surface area of two repeated experiments is about 39.8 and about 41.7 m²/g, respectively.

The conditions are repeated, except instead of directly calcining the wet cake at about 400°C, a pre-drying stage is carried out at about 120°C for about 1 hour. The surface area measured after calcination at about 900°C for about 6 hours is about 36.6 m²/g.

Example 2

Example 1 is repeated but the washing solvent is an alkoxyated compound sold under the designation Dowanol TBH by Dow Chemical Corp. containing an n-butanol chain with a 3 ethoxy group (3EO) or higher. After filtration, the wet cake is washed with about 375 ml of water and then with about 375 ml of a 20% compound solution comprising 20 wt% of the alkoxyated compound and 80 wt. % water. The surface area after calcination at about 900°C for about 6 hours is about 51.7 m²/g.

Example 3

The conditions of Example 1 are repeated but another alkoxyated compound is used comprising a 2 ethylhexyl alcohol with 3EO sold under the designation Rhodasurf by Rhône-Poulenc Inc. The wet cake is impregnated with about 375 ml of deionized water and about 375 ml of a solution comprising 20 wt. % of the compound and 80 wt. % water. The surface area after calcination at about 900°C for about 6 hours is about 47.6 m²/g.

Example 4

In this Example, three alkoxyated compounds are compared with various EO contents: Dowanol, Dowanol DB and Dowanol TBH, n-butanol containing, respectively, 1, 2, and 3 and higher EO. The wet cake is prepared as in Example 1 and is washed with about 375 ml of pure water and then with about 375

ml of a 20 wt % solution of compound. The surface areas after calcination at about 900°C for about 6 hours for the three products are about 45.2, about 49.0 and about 51.7 m²/g, respectively, for the compound containing 1, 2, and 3 and higher EO. The efficiency of the compound washing is correlated to the number of EO groups per organic chain. Thus, even better results can be achieved using molecules containing more than 3 ethoxy groups.

In the following examples, the oxygen buffer ability of mixed oxides produced by different methods are assessed by evaluating oxygen storage capacity in an oxidizing medium and oxygen release capacity in a reducing medium. The test assesses the ability of the product to be oxidized in contact with pulses of oxygen and to be reduced in contact with pulses of carbon monoxide. The carrier gas is pure helium at 10 l/h. Injections of gas are done by 16 ml volume pulses. Pulses of CO are carried out with 5% CO diluted in helium, while pulses of O₂ are carried out by a mixture of 2.5% O₂ in Helium. The gas analysis is performed by chromatography using a thermal conductivity detector. The amount of consumed oxygen leads to the oxygen storage capacity value. The specific value of oxygen storage capacity is expressed as ml of oxygen per gram of material used and is measured at about 400°C.

Example 5

Using a co-precipitation method from a nitrate solution of cerium and zirconium, a mixed oxide is prepared of the composition: 80 wt.% CeO₂ and 20 wt.% ZrO₂. The prepared nitrate solution comprises:

	Ce(NO ₃) ₃	30.31 % CeO ₂ :	2309.47 g,
	ZrO(NO ₃) ₂	14.3 % ZrO ₂ :	699.30 g,
25	H ₂ O ₂	31.75 % H ₂ O ₂ :	435.75 g, and
	Deionized Water:		5778.98 g;

with the components being mixed well until homogenous.

The prepared ammonia solution comprises:

	Ammonia Solution	29.8 % NH ₃ :	947.96 g, and
30	Deionized Water:		6876.00 g;

with the components being mixed well until homogenous.

The nitrate solution is added to the ammonia solution to precipitate out the hydroxides. The pH of the reaction is maintained between about 7 and about 11 and the temperature is room temperature. By reacting the salt mixture and adding ammonia, the mixed hydroxides, corresponding to about 30 g of dried rare earth oxides, are precipitated out from the solution and filtered on a Buchner filter. The cake is washed with about 750 ml of deionized water, then calcined at about 400°C for about 1 hour. The thermal stability of the product is evaluated after calcination in a muffle furnace at about 900°C for about 6 hours and the surface area is measured. The surface area of the aged product is about 33 m²/g and the oxygen storage capacity is about 1.75 ml O₂/g.

Example 6

The experiment described in Example 5 is repeated, except, instead of just washing the cake with water, an additional impregnating stage is included. After water washing, the cake (about 30 g as Rare Earth oxides) is impregnated with about 40 g, of a Dowanol TBH ethoxylated butanol and then calcined at about 600°C for about 2 hours. The thermal stability of the product is evaluated after calcination in a muffle furnace at about 900°C for about 6 hours and the surface area is measured. The surface area of the aged product is about 42 m²/g and the oxygen storage capacity is about 2.85 ml O₂/g.

Example 7

In this example, we prepare a mixed oxide with the following composition Ce_{0.5}Zr_{0.5}O₂ from the co-thermohydrolysis method. In a 25-L glass autoclave (Dietrich) mix the stoichiometric amount of (a) a Ce(IV) nitrate solution with a free acidity of 0.62N and (b) a solution of zirconium nitrate. The total oxide concentration in the reactor is about 80 g/l. The mixture is then heated at about 150°C for about 4 hours in the autoclave under constant stirring. A solution of ammonia is then added to the obtained suspension to raise the pH to 9.5. The pH adjusted solution is stirred for about 30 minutes. The precipitate is separated from

the mother liquor by settling and then re-slurried into water. The suspension is heated at about 100°C for about 2 hours. The mixed oxides are filtrated.

In one case, the wet cake is washed with water only (about 375 ml for about 30 g of dried Rare Earth oxides). The surface area of the product calcined
5 for about 6 hours at about 900°C is about 34 m²/g.

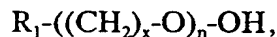
In another case, the wet cake is washed with water under the conditions described before and impregnated with about 40 g of surfactant (Igepal CO-630). The surface area of the product calcined for about 6 hours at about 900°C is about 48 m²/g.

10 Compositions of the present invention can be utilized by known techniques in the art for catalysis of exhaust gas systems and more particularly in catalytic converters for automobiles.

While specific embodiments of the invention have been described in detail, it will be appreciated by those skilled in the art that various modifications and
15 alternatives to those details could be developed in light of the overall teachings of the disclosure. Accordingly, the particular arrangements disclosed are meant to be illustrative only and not limiting as to the scope of invention which is to be given the full breadth of the claims appended and any and all equivalents thereof.

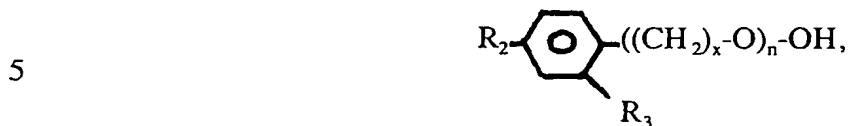
20 What is Claimed is:

1. A composition comprising cerium and zirconium oxides, mixed oxides or solid solutions produced by a method comprising the steps of:
- a) forming a mixture comprising cerium and zirconium mixed oxides, cerium oxides, zirconium oxides, cerium and zirconium solid solutions or mixtures thereof;
 - b) washing or impregnating the mixture with at least one alkoxyated compound having greater than 2 carbon atoms; and
 - c) calcining the washed mixture.
2. The composition of claim 1, wherein the mixture is formed by co-thermohydrolysis or co-precipitation.
3. The composition of claim 1, wherein the mixed oxides have a weight ratio of CeO_2 to ZrO_2 of from about 100:0 to about 5:95.
4. The composition of claim 1, wherein the mixed oxides have a surface area measured after calcination at about 900°C for about six hours of greater than about $50 \text{ m}^2/\text{g}$.
5. The composition of claim 1, wherein the mixed oxides have an oxygen storage capacity of greater than about $2.5 \text{ ml O}_2/\text{g}$, preferably greater than about $2.6 \text{ ml O}_2/\text{g}$.
6. The composition of claim 1, wherein the alkoxyated compound is of the formula:



- wherein R_1 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons and fatty hydrocarbon residues having from 8 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

7. The composition of claim 1, wherein the alkoxyated compound is of the formula:



wherein R₂ and R₃ are the same or different and are independently selected from the group consisting of hydrogen and linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

8. The composition of claim 1, wherein the alkoxyated compound is of the formula:



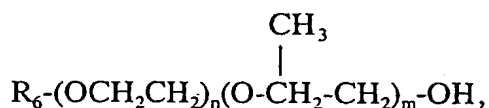
wherein R₄ is selected from the group consisting of linear and nonlinear alcohol groups having from 1 to 20 carbons, preferably R₄ is 2-ethylhexyl alcohol; n is from 1 to 100, preferably n is from 12 to 40 and more preferably n is 3; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

9. The composition of claim 1, wherein the alkoxyated compound is of the formula:



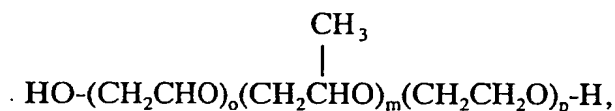
wherein R₅ is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, preferably n is from 12 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

10. The composition of claim 1, wherein the alkoxyated compound is of the formula:



wherein R_6 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and m is from 0 to 200, preferably m is from 0 to 100.

11. The composition of claim 1, wherein the alkoxyated
5 compound is of the formula:



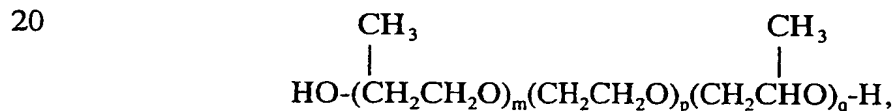
- 10 wherein o is from 0 to 300, m is from 0 to 300, and p is from 0 to 300.

12. The composition of claim 1, wherein the alkoxyated compound is of the formula:



- 15 wherein R_7 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

13. The composition of claim 1, wherein the alkoxyated compound is of the formula:



wherein m is from 0 to 300, p is from 0 to 300, and q is from 0 to 300.

- 25 14. The composition of claim 1, wherein the alkoxyated compound is provided in an aqueous solution and the alkoxyated compound comprises less than about 50, preferably from about 10 to about 30, weight percent of the aqueous solution.

15. The composition of claim 1, wherein the washed mixture is
30 calcined at a temperature of greater than about 300°C.

16. The composition of claim 15, wherein the method for producing the composition further comprises drying the washed mixture at a temperature of less than about 300°C prior to calcining the washed mixture.

17. A method for producing cerium and zirconium oxides, mixed oxides or solid solutions having improved thermal stability, the method comprising the steps of:

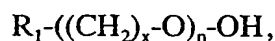
- a) forming a mixture comprising cerium and zirconium mixed oxides, cerium oxides, zirconium oxides, cerium and zirconium solid solution or a mixture thereof;
- b) washing or impregnating the mixture with at least one alkoxyated compound having greater than 2 carbon atoms; and
- c) calcining the washed mixture.

18. The method of claim 17, wherein the mixture is formed by co-thermohydrolysis or co-precipitation.

19. The method of claim 17, wherein the mixture is provided as a wet cake.

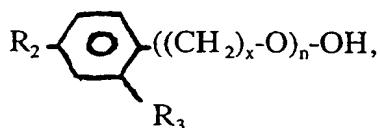
20. The method of claim 17, further comprising:
washing the mixture with water prior to washing or impregnating the mixture with the alkoxyated compound.

21. The method of claim 17, wherein the alkoxyated compound is of the formula:



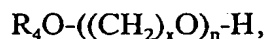
wherein R_1 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons and fatty hydrocarbon residues having from 8 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

22. The method of claim 17, wherein the alkoxyated compound is of the formula:



wherein R_2 and R_3 are the same or different and are independently selected from the group consisting of hydrogen and linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

- 5 23. The method of claim 17, wherein the alkoxyated compound is of the formula:



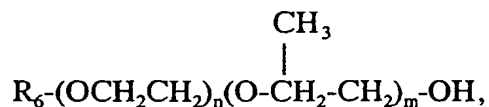
- 10 wherein R_4 is selected from the group consisting of linear and nonlinear alcohol groups having from 1 to 20 carbons, preferably R_4 is 2-ethylhexyl alcohol; n is from 1 to 100, preferably n is 12 to 40 and more preferably n is 3; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

- 15 24. The method of claim 17, wherein the alkoxyated compound is of the formula:



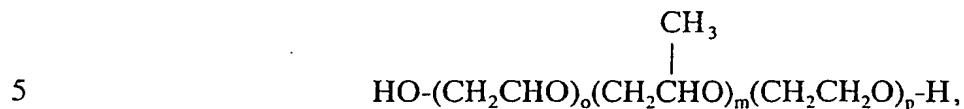
- 20 wherein R_5 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

- 25 25. The method of claim 17, wherein the alkoxyated compound is of the formula:



- 30 wherein R_6 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and m is from 0 to 200, preferably m is from 0 to 100.

26. The method of claim 17, wherein the alkoxyated compound is of the formula:



wherein o is from 0 to 300, m is from 0 to 300, and p is from 0 to 300.

27. The method of claim 17, wherein the alkoxyated compound is of the formula:



wherein R₇ is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

28. The method of claim 17, wherein the alkoxyated compound is of the formula:



wherein m is from 0 to 300, p is from 0 to 300, and q is from 0 to 300.

29. The method of claim 17, wherein the alkoxyated compound is provided in an aqueous solution and the alkoxyated compound comprises less than about 50, preferably from about 10 to about 30, weight percent of the aqueous solution.

30. The method of claim 17, wherein the washed mixture is calcined at a temperature of greater than about 300°C.

31. The method of claim 30, further comprising drying the washed mixture at a temperature of less than about 300°C prior to calcining the washed mixture.

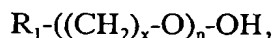
32. The method of claim 17, wherein the calcined mixture has a weight ratio of CeO₂ to ZrO₂ of from about 100:0 to about 5:95.

33. The method of claim 17, wherein the surface area is measured after calcination at about 900°C for about six (6) hours and is greater than about 50 m²/g.

34. The method of claim 17, wherein the calcined mixture has an oxygen storage capacity of greater than about 2.5 ml O₂/g.

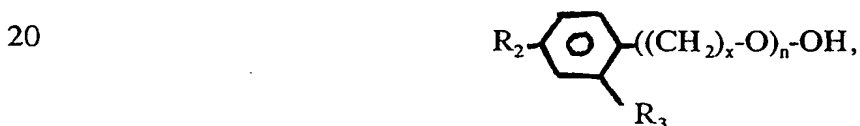
35. A method for improving the thermal stability of cerium and zirconium mixed oxides, cerium oxides, zirconium oxides or (Ce, Zr)O₂ solid solutions, comprising the addition of an alkoxyated compound having greater than 2 carbon atoms to the washing or impregnating step of a co-precipitation or thermohydrolysis process for making said compounds.

36. The method of claim 35 wherein the alkoxyated compound is of the formula:



wherein R₁ is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons and fatty hydrocarbon residues having from 8 to 20 carbons, n is from 1 to 100, and x is from 1 to 4.

37. The method of claim 35, wherein the alkoxyated compound is of the formula:



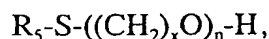
wherein R₂ and R₃ are the same or different and are independently selected from the group consisting of hydrogen and linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4.

38. The method of claim 35, wherein the alkoxyated compound is of the formula:



wherein R₄ is selected from the group consisting of linear and nonlinear alcohol groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4.

39. The method of claim 35, wherein the alkoxyated compound is of the formula:

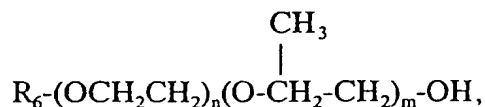


5

wherein R_5 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4.

40. The method of claim 35, wherein the alkoxyated compound is of the formula:

10

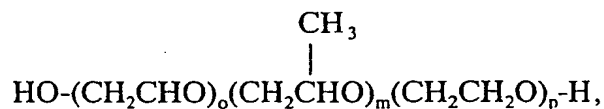


wherein R_6 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and m is from 0 to 200.

15

41. The method of claim 35, wherein the alkoxyated compound is of the formula:

20



wherein o is from 0 to 300, m is from 0 to 300, and p is from 0 to 300.

42. The method of claim 35, wherein the alkoxyated compound is of the formula:

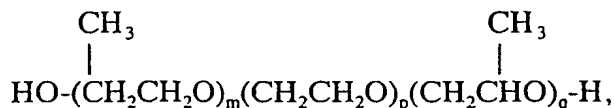
25



wherein R_7 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons, n is from 1 to 100, and x is from 1 to 4.

43. The method of claim 35, wherein the alkoxyated compound is of the formula:

30



wherein m is from 0 to 300, p is from 0 to 300, and q is from 0 to 300.

35

44. The method of claim 35, wherein the alkoxyated compound is added in an aqueous solution and the alkoxyated compound comprises less than about 50, preferably from about 10 to about 30, weight percent of the aqueous solution.

5 45. A method of catalysis for exhaust gas systems comprising utilization of a product of the process comprising the steps of:

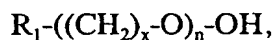
- a) forming a mixture comprising cerium oxides, zirconium oxides, cerium and zirconium mixed oxides, cerium and zirconium solid solution or a mixture thereof;
- 10 b) washing or impregnating the mixture with at least one alkoxyated compound having greater than 2 carbon atoms; and
- c) calcining the washed mixture.

15 46. The method of claim 45, wherein the mixture is formed by co-thermohydrolysis or co-precipitation.

47. The method of claim 45, wherein the mixture is provided as a wet cake.

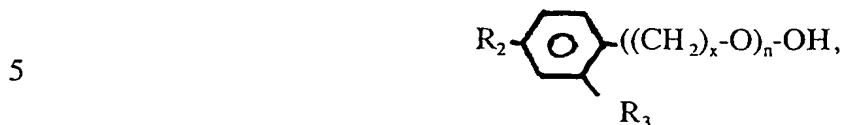
48. The method of claim 45, further comprising:
washing the mixture with water prior to washing or
20 impregnating the mixture with the alkoxyated compound.

49. The method of claim 45, wherein the alkoxyated compound is of the formula:



25 wherein R_1 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons and fatty hydrocarbon residues having from 8 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

50. The method of claim 45, wherein the alkoxyated compound is of the formula:



wherein R_2 and R_3 are the same or different and are independently selected from the group consisting of hydrogen and linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 4 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

51. The method of claim 45, wherein the alkoxyated compound is of the formula:



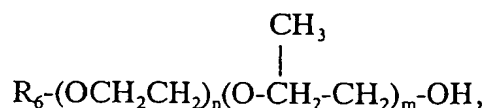
wherein R_4 is selected from the group consisting of linear and nonlinear alcohol groups having from 1 to 20 carbons, preferably R_4 is 2-ethylhexyl alcohol; n is from 1 to 100, preferably n is from 12 to 40 and more preferably n is 3; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

52. The method of claim 45, wherein the alkoxyated compound is of the formula:



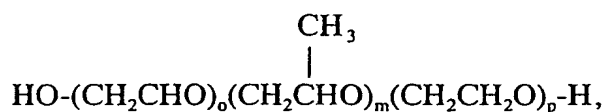
wherein R_5 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

53. The method of claim 45, wherein the alkoxyated compound is of the formula:



wherein R_6 is selected from the group consisting of linear and nonlinear alkyl groups having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and m is from 0 to 200, preferably m is from 0 to 100.

54. The method of claim 45, wherein the alkoxyated compound
5 is of the formula:



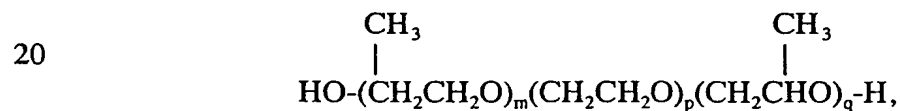
wherein o is from 0 to 300, m is from 0 to 300, and p is from 0 to 300.

55. The method of claim 45, wherein the alkoxyated compound
10 is of the formula:



- wherein R_7 is selected from the group consisting of linear and nonlinear alkyl groups
15 having from 1 to 20 carbons; n is from 1 to 100, preferably n is from 12 to 40; and
 x is from 1 to 4, preferably x is from 1 to 3 and more preferably x is 2.

56. The method of claim 45, wherein the alkoxyated compound
is of the formula:



wherein m is from 0 to 300, p is from 0 to 300, and q is from 0 to 300.

57. The method of claim 45, wherein the alkoxyated compound
is provided in an aqueous solution and the alkoxyated compound comprises less
25 than about 50, preferably from about 10 to about 30, weight percent of the aqueous
solution.

58. The method of claim 45, wherein the washed mixture is
calcined at a temperature of greater than about 300°C.

59. The method of claim 57, further comprising drying the
30 washed mixture at a temperature of less than about 300°C prior to calcining the
washed mixture.

60. The method of claim 45, wherein the calcined mixture has a
weight ratio of CeO_2 to ZrO_2 of from about 100:0 to about 5:95.

61. The method of claim 45, wherein the surface area is measured after calcination at about 900°C for about six (6) hours and is greater than about 50 m²/g.

62. The method of claim 45, wherein the calcined mixture has an
5 oxygen storage capacity of greater than about 2.5 ml O₂/g.

63. A catalytic converter comprising the product of the process comprising the steps of:

- 10
- a) forming a mixture comprising cerium oxides, zirconium oxides, cerium and zirconium mixed oxides, cerium and zirconium solid solution or a mixture thereof;
 - b) washing or impregnating the mixture with at least one alkoxylated compound having greater than 2 carbon atoms; and
 - c) calcining the washed mixture.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/IB 97/01253

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C01G25/00 C01G25/02 B01J23/10 C01B13/14

According to International Patent Classification(IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C01G B01J C01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 391 448 A (NIPPON CATALYTIC CHEM IND) 10 October 1990 see the whole document ---	1,2,6, 14,15
A	EP 0 605 274 A (RHONE POULENC CHIMIE) 6 July 1994 see the whole document ---	1-4
A	EP 0 234 647 A (SOLVAY) 2 September 1987 see the whole document ---	1
A	EP 0 319 365 A (RHONE POULENC CHIMIE) 7 June 1989 -----	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
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- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

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- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

19 January 1998

Date of mailing of the international search report

26/01/1998

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IB 97/01253

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